=> d ibib abs hitstr 1-96

STM-STRUTURE SEARCH

ANSWER 1 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:780372 CAPLUS

TITLE:

Non-steroidal anti-inflammatory drug dosing regimen Gelotte, Cathy K.; Hough, Douglas R.; McNally, Gerard

Ρ.

PATENT ASSIGNEE(S):

USA

SOURCE:

U.S. Pat. Appl. Publ., 11 pp.

CODEN: USXXCO

DOCUMENT TYPE:

INVENTOR(S):

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
·				
US 2004186180	A1	20040923	US 2003-393755	20030321
PRIORITY APPLN. INFO.:			US 2003-393755	20030321
AB A method of admin	istering	non-steroid	lal-anti-inflammatory	drugs, in

particular propionic acid derivs. such as ibuprofen, or acetaminophen is provided. This method provides improved therapeutic effect, in particular pain relief, over extended time periods.

55843-86-2, Miroprofen

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(non-steroidal anti-inflammatory drug dosing regimen)

RN 55843-86-2 CAPLUS

Benzeneacetic acid, 4-imidazo[1,2-a]pyridin-2-yl- α -methyl- (9CI) CN (CA INDEX NAME)

ANSWER 2 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:696342 CAPLUS

DOCUMENT NUMBER:

141:225302

TITLE:

Preparation of N-arylheterocycles as melanin

concentrating hormone (MCH) antagonists.

INVENTOR(S):

Schwink, Lothar; Stengelin, Siegfried; Gossel, Matthias; Boehme, Thomas; Hessler, Gerhard; Stahl,

Petra; Gretzke, Dirk

PATENT ASSIGNEE(S):

Aventis Pharma Deutschland GmbH, Germany

PCT Int. Appl., 390 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

SOURCE:

German

FAMILY ACC. NUM. COUNT:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004072025	A2	20040826	WO 2004-EP1342	20040213
W: AE, AE, AG	, AL, AL	, AM, AM,	AM, AT, AT, AU, AZ, AZ,	BA, BB, BG,
BG. BR. BR	. BW. BY	BY BZ	BZ, CA, CH, CN, CN, CO,	CO. CR. CR.

KG, KZ, MD, RU

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,

GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.:

DE 2002-10237423 A 20020816

AB The invention discloses a method for treatment of immunol. diseases or pathol. conditions which contain an immunol. component, using certain LCK inhibitors, which already are known as kinase inhibitors for therapy in oncol., optionally in combination with one or more other medications selected from NSAIDs, steroids, DMARDs, immunosuppressants, biol. response modifiers, and antiinfectives. Also disclosed are pharmaceutical compns. which contain the LCK inhibitors as well as the other medications, and use of LCK inhibitors for production of a pharmaceutical composition for treatment

o£

immunol. diseases or pathol. conditions which contain an immunol. component.

IT **55843-86-2**, Miroprofen

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(LCK inhibitors for treatment of immunol. diseases, and use with other agents)

RN 55843-86-2 CAPLUS

CN Benzeneacetic acid, 4-imidazo[1,2-a]pyridin-2-yl- α -methyl- (9CI) (CA INDEX NAME)

L4 ANSWER 10 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:101449 CAPLUS

DOCUMENT NUMBER:

140:128417

TITLE:

A process for the synthesis of novel 2-substituted

imidazo[1,2-a]pyridines

INVENTOR(S):

Raj, Kanwal; Vishnoi, Surendra Pal; Shoeb, Aboo; Gupta, Deoki Nandan; Keshri, Govind; Kamboj, Ved

Prakash

PATENT ASSIGNEE(S):

Council of Scientific and Industrial Research, India

SOURCE:

Indian, 8 pp.

CODEN: INXXAP

DOCUMENT TYPE: LANGUAGE: Patent

DANTER AGG NUM COUNTY

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
IN 179790	A	19971206	IN 1992-DE104	19920210
PRIORITY APPLN. INFO.:			IN 1992-DE104	19920210
OTHER SOURCE(S):	CASREA	CT 140:12841	.7; MARPAT 140:128417	
CT				

GI

The title compds. [I; Rl = alkoxy such as 3-ethoxy, dialkylaminoethoxy such as diethylaminoethoxy, cyclic aminoethoxy such as pyrrolidinoethoxy], useful as antiimplantation/abortifacient agents, were prepared by reacting the phenolic compound II with alkyl halides in presence of organic solvent like acetone and K2CO3 at reflux temperature Thus, refluxing 2-(4'-hydroxyphenyl)imidazo[1,2-a]pyridine with N-(2-chloroethyl)pyrrolidine.HCl in the presence of K2CO3 in Me2CO afforded 50% I [Rl = 4-(2-pyrrolidinoethoxy)]. The compds. I showed 30-100% protection against pregnancy at 2.5-20 mg/kg by oral and s.c. routes when tested in hamsters.

651042-78-3P 651042-79-4P 651042-80-7P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
(Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(a process for the synthesis of novel 2-substituted imidazo[1,2-a]pyridines as antifertility agents)

RN 651042-78-3 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-[4-[2-(1-pyrrolidinyl)ethoxy]phenyl]- (9CI) (CA INDEX NAME)

RN 651042-79-4 CAPLUS

CN Ethanamine, 2-(4-imidazo[1,2-a]pyridin-2-ylphenoxy)-N,N-dimethyl- (9CI) (CA INDEX NAME)

RN 651042-80-7 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-(3-ethoxyphenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

L4 ANSWER 16 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:14078 CAPLUS

DOCUMENT NUMBER:

140:59640

TITLE:

A process for the preparation of 2-substituted and 2,5,7-tri-substituted imidazo(1,2a)pyridines via cyclocondensation reaction showing antifertility

activity

INVENTOR(S):

Raj, Kanwal; Vishnoi, Surendra Pal; Gupta, Deoki Nandan; Keshri, Govind; Kamboj, Ved Prakash

PATENT ASSIGNEE(S):

Council of Scientific and Industrial Research, India

SOURCE:

Indian, 11 pp.

CODEN: INXXAP

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

. 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
IN 179788 PRIORITY APPLN. INFO.:	A	19971206	IN 1992-DE102 IN 1992-DE102	19920210 19920210
OTHER SOURCE(S):	CASRE	ACT 140:59640); MARPAT 140:59640	
GI				-

$$R^{1}$$
 R^{2}
 R^{3}
 R^{3}
 R^{2}
 R^{3}
 R^{3

A process for the preparation of novel 2-substituted and 2,5,7 trisubstituted AB imidazo(1,2-a)pyridines I, where R1 represents substituted phenol such as halophenyl preferably chloro or bromo Ph, Me-Ph, nitro-Ph, hydroxy-Ph, methoxy-Ph, ethoxy-Ph, carboxymethyl, carbethoxymethyl; R2= R3 = H, CH3, which comprises refluxing correspondingly substituted 2-aminopyridines of formula II where R2 and R3 represent hydrogen or Me groups with correspondingly substituted - bromoketone of the formula III where R1 represents substituted phenol such as halophenyl preferably chloro or bromophenyl, alkyl, Ph, nitro Ph, hydroxy Ph, methoxy Ph, ethoxy Ph or carboxymethyl and carbethoxymethyl; in the presence of an aprotic solvent for a period ranging from 2-24 h. at a temperature in the range $36^{\circ}-90^{\circ}\text{C}$ and recovering the 2-substituted and 2,5,7 trisubstituted imidazo(1,2-a)pyridines, of formula I by conventional methods. Thus, 2-(4-hydroxyphenyl)imidazo[1,2-a]pyridine hydrochloride was prepared as antifertility agent. The title compds. prepared by the process of the present invention found to be useful as antiimplantation abortifacient agents. These compds. when tested in hamster for antifertility activity showed 30-100% protection against pregnancy at 2.5 - 20 mg/kg by oral and s.c. routes.

IT 88467-84-9P 637768-42-4P

RL: IMF (Industrial manufacture); PAC (Pharmacological activity); SPN (Synthetic preparation); **THU (Therapeutic use)**; BIOL (Biological study); PREP (Preparation); USES (Uses)

(process for preparation of substituted and substituted imidazoapyridines via cyclocondensation reaction showing antifertility activity)

RN 88467-84-9 CAPLUS

Phenol, 2-imidazo[1,2-a]pyridin-2-yl-, monohydrobromide (9CI) (CA INDEX CN NAME)

HBr

637768-42-4 CAPLUS RN

Phenol, 4-imidazo[1,2-a]pyridin-2-yl-, monohydrobromide (9CI) (CA INDEX CNNAME)

HBr

ANSWER 17 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:2623 CAPLUS

DOCUMENT NUMBER:

140:59409

TITLE:

Preparation of hydroxamate derivatives of nonsteroidal

antiinflammatory drugs

INVENTOR(S):

Wang, Tingmin; Lai, Ching-San

PATENT ASSIGNEE(S):

Medinox, Inc., USA PCT Int. Appl., 58 pp.

SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

3

PATEN	1 T	10.			KINI	O I	DATE		1	APPL	I CAT:	ION I	NO.		DA	ATE	
WO 20					A2 A3		2003 2004		Ţ	WO 2	003-T	JS19:	228		20	0030	617
ħ	N:	CO, GM, LS, PG,	CR, HR, LT, PH,	CU, HU, LU, PL, UA,	CZ, ID, LV, PT, UG,	DE, IL, MA, RO,	DK, IN, MD, RU,	DM, IS, MG, SC,	DZ, JP, MK, SD,	EC, KE, MN, SE,	EE, KG, MW, SG,	ES, KP, MX, SK,	FI, KR, MZ, SL,	GB, KZ, NI, TJ,	CA, GD, LC, NO, TM, AZ,	GE, LK, NZ, TN,	GH, LR, OM, TR,
F	₹W:	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	AT, IT, GA,	LU,	MC,

US 6720324

B2 20040413

PRIORITY APPLN. INFO.:

US 2000-216218P

P 20000705

OTHER SOURCE(S):

MARPAT 138:287697

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Title compds. I-IV [A = (un)substituted Ph, pyridyl, benzothiazolyl,AΒ benzoxazolyl, etc.; R1 = H, NO2, CN, (fluoro)alkyl, (cyclo)alkenyl, alkynyl, (fluoro)cycloalkyl, amino, alkoxy, acyl, carboxy, carboxamido; R2 = H, (hydroxy)alkyl, alkoxyalkyl, fluoroalkyl, cycloalkenyl, etc.; R3 = H, (fluoro)alkyl, (cyclo)alkenyl, alkynyl, (fluoro)cycloalkyl; R4 = alkyl-piperidinyl, alkyl-tetrahydropyridinyl, etc. in which the heterocycle is substituted with (hetero)aryl, thioacyl, amido, etc.; X = O, S, NR3; n = 0 - 5] were prepared For instance, (+)-V was prepared by reaction of 5-methoxycarbonyl-4-methoxymethyl-1,2,3,6-tetrahydro-2-oxo-6-(3,4-difluorophenyl)-1-[(4-nitrophenyloxy)carbonyl]pyrimidine (preparation given) and the corresponding propylamine sidechain with base (e.g., iPr2NEt) in CH2Cl2. (+)-V had antagonist potency (Kb) = 0.3 nM and Ki = 0.08 nM for the melanin-concentrating hormone receptor (mch) and Ki > 50,000 nM for two neuropeptide Y receptors and Ki > 50,000 nM three galanin receptors. I-IV are useful in the treatment of, e.g., bulimia nervosa and obesity.

IT 330963-56-9P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); **THU** (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug; preparation and use of arylpyrimidines as selective melanin concentrating

hormone-1 (mch-1) receptor antagonists)

RN 330963-56-9 CAPLUS

CN 1,3-Benzenediol, 4-[[(4-imidazo[1,2-a]pyridin-2-ylphenyl)imino]methyl]-5-methyl- (9CI) (CA INDEX NAME)

ANSWER 40 OF 96

CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:888557 CAPLUS

DOCUMENT NUMBER:

137:384841

TITLE:

Preparation of imidazo[1,2-a]pyridines as mGluR5

antagonists

INVENTOR(S):

Mutel, Vincent; Peters, Jens-Uwe; Wichmann, Juergen

F. Hoffmann-La Roche A.-G., Switz.

PATENT ASSIGNEE(S): SOURCE:

PCT Int. Appl., 31 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

```
APPLICATION NO.
                                                                   DATE
                         KIND
                                DATE
    PATENT NO.
                                            ______
                                                                   ______
                                                                   20020320
                                            WO 2002-EP3098
    WO 2002092086
                         Α1
                                20021121
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL,
            PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,
            UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH. GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
            CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
            BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                            US 2002-93790
                                                                   20020308
    US 2002188128
                          Α1
                                20021212
                          B2
                                20030722
    US 6596731
                                            EP 2002-737889
                                                                   20020320
                                20040121
    EP 1381363
                         Α1
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                         Α
                                20040615
                                            BR 2002-8387
                                                                   20020320
    BR 2002008387
                                            JP 2002-589003
                                                                   20020320
    JP 2004525192
                         T2
                                20040819
    US 2003212096
                         A1
                                20031113
                                            US 2003-407928
                                                                   20030404
                                20040916
                                            US 2004-809068
                                                                   20040325
                         Α1
    US 2004180921
                                            EP 2001-107562
                                                                A 20010327
PRIORITY APPLN. INFO.:
                                            US 2002-93790
                                                                A3 20020308
                                                                W
                                                                   20020320
                                            WO 2002-EP3098
                                            US 2003-407928
                                                                A3 20030404
OTHER SOURCE(S):
                         MARPAT 137:384841
```

GΙ

 R^{1}

$$\mathbb{R}^1$$
 \mathbb{N} \mathbb{A} \mathbb{R}^2

The title compds. [I; R1, R2 = H, alkyl, halogen, OH, alkoxy; A =AB(un) substituted aryl, heteroaryl, etc.], useful for the manufacture of medicaments for the treatment or prevention of GluR5 receptor mediated disorders, such as acute and/or chronic neurol. disorders, were prepared and formulated. Thus, reacting 2-amino-4-chloropyridine with 3,4-dimethylphenacyl bromide in EtOH afforded 69% 7-chloro-2-(3,4dimethylphenyl)imidazo[1,2-a]pyridine which showed IC50 of 0.1 µM against mGluR 5a receptor binding.

```
IT
    885-91-6P 34658-67-8P 65964-60-5P
    88965-00-8P 100965-76-2P 158959-20-7P
    205655-15-8P 326018-20-6P 419557-33-8P
     420832-11-7P 475992-33-7P 475992-34-8P
     475992-35-9P 475992-36-0P 475992-37-1P
     475992-38-2P 475992-39-3P 475992-40-6P
     475992-41-7P 475992-42-8P 475992-43-9P
     475992-44-0P 475992-45-1P 475992-46-2P
     475992-47-3P 475992-48-4P 475992-49-5P
     475992-50-8P 475992-51-9P
```

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES

(preparation of imidazo[1,2-a]pyridines as mGluR5 antagonists)

RN885-91-6 CAPLUS

Imidazo[1,2-a]pyridine, 7-methyl-2-phenyl- (7CI, 8CI, 9CI) CNNAME)

RN 34658-67-8 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-(3-nitrophenyl)- (9CI) (CA INDEX NAME)

RN 65964-60-5 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-(4-methylphenyl)- (9CI) (CA INDEX NAME)

RN 88965-00-8 CAPLUS

CN Imidazo[1,2-a]pyridine, 6-methyl-2-(4-methylphenyl)- (9CI) (CA INDEX NAME)

RN 100965-76-2 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-(2-benzofuranyl)- (6CI, 9CI) (CA INDEX NAME)

RN 158959-20-7 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-(3-chlorophenyl)- (9CI) (CA INDEX NAME)

RN 205655-15-8 CAPLUS

PAGE 2-A

REFERENCE COUNT:

THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 44 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:772124 CAPLUS

DOCUMENT NUMBER:

137:279190

TITLE:

Preparation of imidazo[1,2-a]pyridines as NO synthase

inhibitors

INVENTOR(S):

Maul, Corinna; Hennies, Hagen-Heinrich; Sundermann,

Bernd

PATENT ASSIGNEE(S):

Gruenenthal G.m.b.H., Germany

SOURCE:

Ger. Offen., 12 pp. CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10117183	A1	20021010	DE 2001-10117183	20010405
WO 2002080914	A2	20021017	WO 2002-EP3795	20020405
WO 2002080914	A3	20030103		
W. AF AC AL	ΔΜ ΔΤ	ΔΙΙ. Δ7 ΒΔ	RR RC RP RV R7	CV CH CM

```
CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM,
             HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
             LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL,
             PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA,
             UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
             CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     EP 1372647
                          A2
                                20040102
                                          EP 2002-727529
                                                                    20020405
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
     JP 2004529141
                          T2
                                20040924
                                            JP 2002-578953
                                                                    20020405
     US 2004142961
                          Α1
                                20040722
                                            US 2003-678645
                                                                    20031003
PRIORITY APPLN. INFO.:
                                            DE 2001-10117183
                                                                    20010405
                                                                 Α
                                            WO 2002-EP3795
                                                                    20020405
                                                                 W
                         MARPAT 137:279190
OTHER SOURCE(S):
```

 R^2

R3

Ι

The title compds. [I; R1, R2 = alkyl, cycloalkyl, cycloalkylalkylene, (substituted) aryl, heteroaryl, H, F, Cl, Br, I, cyano, NO2, amino, COR5, CO2H, CO2R6, OH, OR7; R3 = alkyl, cycloalkyl, cycloalkylalkylene, (substituted) aryl, heteroaryl, arylalkylene, heteroarylalkylene, CH2SR8, CH2OR8, H; R4 = H, alkyl, cycloalkyl, heterocyclyl, (substituted) aryl, heteroaryl, etc.; R5 = alkyl, cycloalkyl, cycloalkylalkylene, heterocyclyl, (substituted) aryl, heteroaryl, arylalkylene, heteroarylalkylene; R6, R7 = alkyl, cycloalkyl, cycloalkylalkylene, (substituted) aryl, heteroaryl, arylalkylene, heteroarylalkylene; R8 = alkyl, (substituted) aryl, heteroaryl, arylalkylene, heteroarylalkylene, cycloalkyl], were prepared Thus, 2-bromo-4-methoxyacetophenone was added to 2-amino-4-methylpyridine in EtOH followed by reflux for 2 h and stirring over night at room temperature to give 69% 2-(4-methoxyphenyl)-7-methylimidazo[1,2-a]pyridine. The latter at 10 μM inhibited NO

IT 65964-63-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); **THU** (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of imidazopyridines as NO synthase inhibitors)

RN 65964-63-8 CAPLUS

synthase by 39%.

CN Imidazo[1,2-a]pyridine, 2-(4-methoxyphenyl)-7-methyl- (9CI) (CA INDEX NAME)

7

REFERENCE COUNT:

study, unclassified); THU (Therapeutic use); BIOL (Biological

study); USES (Uses)

(peroral meloxicam suspension)

55843-86-2 CAPLUS RN

Benzeneacetic acid, 4-imidazo[1,2-a]pyridin-2-yl- α -methyl- (9CI) CN

(CA INDEX NAME)

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 4 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 60 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1999:454955 CAPLUS

DOCUMENT NUMBER:

131:110800

TITLE:

Determination of non-steroidal anti-inflammatory drugs

in biological fluids

AUTHOR(S):

Hercegova, Andrea; Polonsky, J.

CORPORATE SOURCE:

Department Analytical Chemistry, Faculty Chemical Technology, Slovak Univ. Technology, Bratislava,

81237, Slovakia

SOURCE:

Pharmazie (1999), 54(7), 479-486

CODEN: PHARAT; ISSN: 0031-7144 Govi-Verlag Pharmazeutischer Verlag

PUBLISHER:

DOCUMENT TYPE:

Journal; General Review

LANGUAGE: English

A review with 73 refs. is given on HPLC, high performance thin-layer AB chromatog. (HPTLC), liquid chromatog. (LC), GC/MS, capillary electrophoresis, and optical methods for the determination of non-steroidal

anti-inflammatory drugs like arylpropionic acids in urine, serum, and

plasma.

55843-86-2, Miroprofen TT

RL: ANT (Analyte); THU (Therapeutic use); ANST (Analytical

study); BIOL (Biological study); USES (Uses)

(determination of non-steroidal anti-inflammatory drugs in biol. fluids)

RN 55843-86-2 CAPLUS

CN Benzeneacetic acid, 4-imidazo[1,2-a]pyridin-2-yl- α -methyl- (9CI) (CA INDEX NAME)

73 THERE ARE 73 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 61 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1999:271040 CAPLUS

DOCUMENT NUMBER:

130:347408

TITLE: Imidazole[1,2-a]pyridine derivatives as inhibitors for

STAT6 6 transcription factor activation

Inoue, Tadahiro; Iwai, Kiyotaka; Murata, Masashi; INVENTOR(S):

Nishinaka, Shiqeyuki; Aoki, Mikio; Kawakami, Hajime

Sumitomo Pharmaceuticals Co., Ltd., Japan PATENT ASSIGNEE(S):

Jpn. Kokai Tokkyo Koho, 14 pp. SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

FAMILY ACC. NUM. COUNT:

Japanese

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 11116481 PRIORITY APPLN. INFO.:	A2	19990427	JP 1997-290258 JP 1997-290258	19971006 19971006

OTHER SOURCE(S): MARPAT 130:347408

Imidazole[1,2-a]pyridine derivs. (Markush's structure given) and their pharmaceutically acceptable salts are claimed as inhibitors for STAT6 6 transcription factor activation and IL 4 antagonists for treatment of allergic, autoimmune, parasital, viral, and bacterial diseases, tumors. host-vs. graft syndrome, systemic lupus erythematosus, and AIDS.

158959-18-3P 174629-57-3P 224633-75-4P TΤ 224633-76-5P 224633-77-6P 224633-78-7P 224633-79-8P 224633-80-1P 224633-81-2P 224633-82-3P 224633-83-4P 224633-85-6P 224633-95-8P 224633-99-2P 224634-01-9P

224634-03-1P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (imidazole[1,2-a]pyridine derivs. as inhibitors for STAT6 6

transcription factor activation)

RN 158959-18-3 CAPLUS

CNImidazo[1,2-a]pyridine, 2-(3,4-dichlorophenyl)- (9CI) (CA INDEX NAME)

RN 174629-57-3 CAPLUS

Imidazo[1,2-a]pyridine, 2-(4-chlorophenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

HBr

CN Imidazo[1,2-a]pyridine, 2-(2-naphthalenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224633-76-5 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-(4-methylphenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224633-77-6 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-(4-bromophenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224633-78-7 CAPLUS

CN Imidazo[1,2-a]pyridine, 6-chloro-2-(4-fluorophenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224633-79-8 CAPLUS

CN Imidazo[1,2-a]pyridine, 6-chloro-2-(4-chlorophenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224633-80-1 CAPLUS

CN Imidazo[1,2-a]pyridine, 6-chloro-2-(3-methoxyphenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \text{Cl} & & \\ \hline & N & \\ \hline & N & \\ \end{array}$$
 OMe

• HBr

RN 224633-81-2 CAPLUS

CN Imidazo[1,2-a]pyridine, 6-chloro-2-(2-naphthalenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

CN Imidazo[1,2-a]pyridine, 2-[1,1'-biphenyl]-4-yl-6-methyl-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224633-83-4 CAPLUS
CN Imidazo[1,2-a]pyridine, 6-methyl-2-(4-methylphenyl)-, monohydrobromide
(9CI) (CA INDEX NAME)

• HBr

RN 224633-85-6 CAPLUS
CN Imidazo[1,2-a]pyridine, 7-methyl-2-(4-methylphenyl)-, monohydrobromide
(9CI) (CA INDEX NAME)

• HBr

• HBr

RN 224633-99-2 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-[4-(trifluoromethyl)phenyl]-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224634-01-9 CAPLUS

CN Imidazo[1,2-a]pyridine, 6-chloro-2-(4-methylphenyl)-, monohydrobromide (9CI) (CA INDEX NAME)

• HBr

RN 224634-03-1 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-[1,1'-biphenyl]-4-yl-, monohydrobromide (9CI) (CA INDEX NAME)

IT 207277-26-7P 207277-27-8P 207277-28-9P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 2-methoxyphenylpiperazine derivs. as antipsychotics)

RN 207277-26-7 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-[4-[2-[4-(2-methoxyphenyl)-1-piperazinyl]ethoxy]phenyl]- (9CI) (CA INDEX NAME)

RN 207277-27-8 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-[4-[3-[4-(2-methoxyphenyl)-1-piperazinyl]propoxy]phenyl]- (9CI) (CA INDEX NAME)

RN 207277-28-9 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-[4-[4-[4-(2-methoxyphenyl)-1-piperazinyl]butoxy]phenyl]- (9CI) (CA INDEX NAME)

REFERENCE COUNT:

THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 67 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

1

ACCESSION NUMBER:

1998:268348 CAPLUS

DOCUMENT NUMBER:

128:321662

TITLE:

Compositions and methods for treating bone deficit

conditions

INVENTOR(S):

Orme, Mark W.; Baindur, Nand; Robbins, Kirk G.; et al.

PATENT ASSIGNEE(S):

Zymogenetics, Inc., USA; Osteoscreen, Inc.

SOURCE: PCT Int. Appl., 215 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

LANGUAGE:

FILE

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.			DATE
WO 9817267 W: AL, AM, AU, KG, KP, KR,	A1 19980430 BB, BG, BR, CA, LK, LR, LT, LV,	WO 1997-US18864 CN, CZ, EE, FI, GE, HU MD, MG, MK, MN, MX, NO US, US, US, US, US, US	, NZ, PL, RO,
RW: GH, KE, LS, GB, GR, IE,	MW, SD, SZ, UG,	KZ, MD, RU, TJ, TM ZW, AT, BE, CH, DE, DK PT, SE, BF, BJ, CF, CG	
US 5990169 US 6153631 US 6251901 US 5919808 US 5922753 US 5948776 US 5994358 US 6342514 US 5965573 AU 9749889 EP 973513	A 19991123 A 20001128 B1 20010626 A 19990706 A 19990713 A 19991130 B1 20020129 A 19991012 A1 19980515 A1 20000126	US 1997-806768 US 1997-806769 US 1997-808743 US 1997-808742 US 1997-808739 US 1997-808744 US 1997-808741 US 1997-812141 AU 1997-49889	19971023
IE, FI	T2 20010731 B1 20031118	JP 1998-519529 US 1999-297188 US 1996-735870 US 1996-735873 US 1996-735874 US 1996-735881 US 1996-736220 US 1996-736221 US 1996-736222 US 1996-736228 US 1996-736318 US 1996-736319	19971023 19991119 A2 19961023 A2 19961023
OTHER SOURCE(S):	MARPAT 128:3216		

$$^{\mathrm{F}_{3}\mathrm{C}}$$
 $^{\mathrm{H}}$
 $^{\mathrm{N}}$
 $^{\mathrm{CF}_{3}}$
 $^{\mathrm{II}}$

AB Compds. containing 2 covalently linked aromatic systems, i.e. ArlLAr2 [I; Arl, Ar2 = (un)substituted Ph, naphthyl, or 5- or 6-membered aromatic heterocyclyl; L = linker (atoms or covalent bond per se) so as to space the aromatic systems at a distance of 1.5-15 Å] are effective in treating conditions associated with bone deficits. The compds. can be administered to vertebrate subjects alone or in combination with addnl. agents that

promote bone growth or that inhibit bone resorption. They can be screened for activity prior to administration by assessing their ability to effect the transcription of a reporter gene coupled to a promoter associated with a bone morphogenetic protein and/or their ability to stimulate calvarial growth in model animal systems. A variety of compds. were prepared and/or tested by high-throughput screening. For instance, title compound II was prepared by condensation of 2-chloro-5-(trifluoromethyl)pyridine with ethylenediamine in the presence of EtN(Pr-iso)2 at reflux. At 5-50 μq/kg/day in ovariectomized rats, II stimulated bone growth with volume increases of 21-71% observed In a calvarial bone growth assay, another compound I induced a 4-fold increase in width of new calvarial bone vs. controls.

IT31562-99-9 205655-12-5

> RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(preparation and/or use of linked aromatic and heteroarom. compds. for treating

bone deficit conditions)

31562-99-9 CAPLUS RM

CN Imidazo[1,2-a]pyridine, 2-(4-methoxyphenyl)- (9CI) (CA INDEX NAME)

205655-12-5 CAPLUS RN

CNPhenol, 3-imidazo[1,2-a]pyridin-2-yl- (9CI) (CA INDEX NAME)

REFERENCE COUNT:

12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 68 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1998:55635 CAPLUS

DOCUMENT NUMBER:

128:114954

TITLE:

Preparation and formulation of thienyloxadiazole derivatives and analogs as anti-phencyclidine agents

INVENTOR (S):

Kimura, Takenori; Murakami, Takeshi; Ohmori, Junya;

Morita, Takuma; Tsukamoto, Shin-ichi

PATENT ASSIGNEE(S):

Yamanouchi Pharmaceutical Co., Ltd., Japan

SOURCE: PCT Int. Appl., 90 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

1

FAMILY ACC. NUM. COUNT:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	-			
WO 9800420	A1	19980108	WO 1997-JP2255	19970630
W: AL, AM, AU,	AZ, BA	BB. BG. BR	BY, CA, CN, CU, CZ,	EE GE GH

RN 144701-75-7 CAPLUS

CN 1H-Benzimidazole, 2-butyl-6-imidazo[1,2-a]pyridin-2-yl-4-methyl-1-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (9CI) (CA INDEX NAME)

L4 ANSWER 86 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1992:83664 CAPLUS

DOCUMENT NUMBER:

116:83664

TITLE:

Preparation of 5,6,7,8-tetrahydro-4H-thiazolo[5,4-

blazepine derivatives as antihypertensives

INVENTOR(S):

Aono, Tetsuya; Shimamoto, Norio

PATENT ASSIGNEE(S):

Takeda Chemical Industries, Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 63 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 03206042	A2	19910909	JP 1990-833	19900106
PRIORITY APPLN. INFO.:			JP 1990-833	19900106
OTHER SOURCE(S):	MARPAT	116:83664		

GI

$$\begin{array}{c|c}
 & N \\
 & N \\
 & R^2
\end{array}$$

The title compds. [I; R1 = H, (un) substituted aliphatic, acyl or sulfonyl; R2= H, (un)substituted aromatic or aliphatic] are prepared as K channel opener. Thus, 14.8 q 1,1'-carbonyldiimidazole was added to a solution of 12 g 2,6-F2C6H3CO2H in THF and thereto after stirring 15 min at room temperature

9.73

CN

q 3-amino-e-caprolactam was added and the mixture was stirred 5 h at room temperature to give 13.5 g 3-(2,6-difluorobenzoylamino)- ϵ caprolactam which (8.96 g) was refluxed 24 h, with 8.96 g P4S10 in pyridine to give 23.8% I (R1 = H, R2 = 2,6-F2C6H3)(II). II and I [R1 = H, R2 = 2,6-F2C6H3](II). R2 = (Z)-4-Et2NC6H4CH:CH] (III) in vitro inhibited 8 and 100%, resp., rat aorta contraction induced by Et3NCl and BaCl2 and gave no inhibition of the one induced by 80 mM KCl. II and III at 1 mg/kg i.v. lowered 49 and 46%, resp. the blood pressure of rats. A total of 175 I were prepared

IT 138621-33-7P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation of, as antihypertensive)

138621-33-7 CAPLUS RN

4H-Thiazolo[5,4-b]azepine, 5,6,7,8-tetrahydro-2-imidazo[1,2-a]pyridin-2-yl-, dihydrochloride (9CI) (CA INDEX NAME)

2 HC1

CAPLUS COPYRIGHT 2004 ACS on STN ANSWER 87 OF 96

Journal

ACCESSION NUMBER: 1988:570365 CAPLUS

DOCUMENT NUMBER: 109:170365

TITLE: Synthesis of (aryloxy)alkylamines. 2. Novel imidazo-fused heterocycles with calcium channel

blocking and local anesthetic activity

AUTHOR (S): Sanfilippo, Pauline J.; Urbanski, Maud; Press, Jeffery

B.; Dubinsky, Barry; Moore, John B., Jr.

CORPORATE SOURCE: Res. Lab., Ortho Pharm. Corp., Raritan, NJ, 08869, USA SOURCE: Journal of Medicinal Chemistry (1988), 31(11), 2221-7

CODEN: JMCMAR; ISSN: 0022-2623

DOCUMENT TYPE:

LANGUAGE: English

OTHER SOURCE(S): CASREACT 109:170365

ANSWER 96 OF 96 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

1975:471745 CAPLUS

DOCUMENT NUMBER:

83:71745

TITLE:

Pharmacology of zolimidine [2-(p-

methylsulfonylphenyl)imidazo(1,2-a)pyridine], a new

nonanticholinergic gastroprotective agent. I.

Protective activity against experimental neurogenic gastric ulcers and other pharmacologic properties

AUTHOR (S):

Murmann, W.; Carminati, G. M.; Cattaneo, R. Res. Dep., Selvi C. S.p.A., Milan, Italy Panminerva Medica (1974), 16(10), 301-20

SOURCE:

CODEN: PMMDAE; ISSN: 0031-0808

DOCUMENT TYPE:

CORPORATE SOURCE:

Journal English

LANGUAGE:

For diagram(s), see printed CA Issue. GT

Zolimidine (I) [1222-57-7] had low toxicity as well as a potent inhibitor AΒ of qastric ulcers of neurogenic origin in rats. At doses slightly higher than antiulcer doses, I exerted antiinflammatory and antipyretic actions; at yet somewhat higher doses it exerted central sedation without affecting reflex activity. Muscle relaxant and antihistaminic activities were also demonstrated in vitro. Both in vitro and in vivo the drug is totally devoid of anticholinergic activity. I appears to exert its action directly on the gastric mucus cells.

IT 1222-57-7

> RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(pharmacol. of)

RN 1222-57-7 CAPLUS

CN Imidazo[1,2-a]pyridine, 2-[4-(methylsulfonyl)phenyl]- (9CI) (CA INDEX NAME)

=> d his

(FILE 'HOME' ENTERED AT 15:32:42 ON 08 OCT 2004)

FILE 'REGISTRY' ENTERED AT 15:32:55 ON 08 OCT 2004

L1STRUCTURE UPLOADED

L218 S L1

L31944 S L1 FULL

FILE 'CAPLUS' ENTERED AT 15:35:02 ON 08 OCT 2004

L4

96 S L3/THU

=> d 11

L1 HAS NO ANSWERS

L1

STR

G1 H,O,X,Ak

Structure attributes must be viewed using STN Express query preparation.

=>



PALM INTRANET

Day: Tuesday Date: 10/12/2004 Time: 11:58:08

Inventor Name Search Result

Your Search was:

Last Name = MUTEL First Name = VINCENT

Application#	Patent#	Status	Date Filed	Title
10809068	Not Issued	030	03/25/2004	SUBSTITUTED IMIDAZO[1,2-A]PYRIDINE DERIVATIVES
10438361	6803381	150	05/14/2003	CARBAMIC ACID DERIVATIVES
10407929	Not Issued	041	04/04/2003	PHENYLETHYNYL AND STYRYL DERIVATIVES OF IMIDAZOLE AND FUSED RING HETEROCYCLES
10407928	Not Issued	094	04/04/2003	SUBSTITUTED IMIDAZO [1,2-A] PYRIDINE DERIVATIVES
<u>10397908</u>	Not Issued	041	03/26/2003	1-SULFONYL PYRROLIDINE DERIVATIVES
10396172	Not Issued	041	03/25/2003	PHENYLETHYNYL AND STYRYL DERIVATIVES OF IMIDAZOLE AND FUSED RING HETEROCYCLES
10300449	Not Issued	080	11/20/2002	GLUTAMATE RECEPTOR ANTAGONISTS
10157338	<u>6673795</u>	150		PYRIMIDINE, PYRAZINE AND TRIAZINE DERIVATIVES
<u>10135150</u>	Not Issued	161	04/29/2002	4-AMINOPYRIMIDINE DERIVATIVES
<u>10116597</u>	<u>6548495</u>	150	04/03/2002	DIHYDRO-BENZO [B] [1,4] DIAZEPIN-2-ONE DERIVATIVES
<u>10115826</u>	6544985	150	04/03/2002	DIHYDRO-BENZO [B] [1,4] DIAZEPIN-2-ONE DERIVATIVES
10093790	<u>6596731</u>	150	03/08/2002	SUBSTITUTED IMIDAZO[1,2-A] PYRIDINE DERIVATIVES
10020680	6586422	150		PYRAZINE AND TRIAZINE DERIVATIVES OF 1,2,4,5-TETRAHYDRO-BENZO OR THIENO [D] AZEPINE
10008827	6596743	150	12/10/2001	CARBAMIC ACID DERIVATIVES
09996641	6706707	150		PHENYLETHYNYL AND STYRYL DERIVATIVES OF IMIDAZOLE AND FUSED RING HETEROCYCLES

<u>09902916</u>	6369222	150	07/11/2001	NOVEL MGLUR ANTAGONISTS AND A METHOD FO THEIR SYNTHESIS
09892994	6399641	150	06/27/2001	NOVEL 2H-TETRAZOLE-AMIDE COMPOUNDS WITH THERAPEUTIC ACTIVITY AS METABOTROPIC GLUTAMATE RECEPTOR AGONISTS
09880539	6589978	150	06/13/2001	1-SULFONYL PYRROLIDINE DERIVATIVES
09687241	6509328	150	10/13/2000	GLUTAMATE RECEPTOR ANTAGONISTS
09687240	6407094	150	10/13/2000	GLUTAMATE RECEPTOR ANTAGONISTS
09669583	6548522	150	09/26/2000	METHOD FOR TREATING CONDITIONS RELATED T THE GLUTAMATE RECEPTOR USING CARBOXYLIC ACID AMIDE DERIVATIVES
09630702	6218385	150	08/01/2000	1,2,4,5-TETRAHYDRO-BENZO[D]AZEPIN DERIVATIVES
09545622	6462198	150	04/10/2000	CARBAMIC ACID DERIVATIVES
09534380	<u>6284785</u>	150	()	1-ARENESULFONYL-2-ARYL-PYROLIDINE AND PYRIDINE DERIVATIVES
09385935	6107342	150	08/30/1999	2-AMINO-BICYCLO[3 1 0]HEXANE-2,6-DICARBOXY ACID DERIVATIVES AND A PROCESS FOR THE PREPARATION THEREOF
09320307	6071929	150	05/26/1999	OCTAHYDRO PHENANTHRIDINE DERIVATES USEF AS NMDA-R RECEPTOR SUBTYPE BLOCKERS
<u>09304624</u>	6054588	150	05/04/1999	HETEROCYCLIC VINYL ETHERS
<u>09304561</u>	Not Issued	161	05/04/1999	HETEROCYCLIC VINYL ETHERS
09121737	Not Issued	161	07/23/1998	HETEROCYCLIC VINYL ETHERS
09115191	5958931	150	07/14/1998	PHENYL-SUBSTITUTED 5H-THIAZOLO[3,2-A] PYRIMIDINE DERIVATIVE GLUTAMATE RECEPTOF ANTAGONISTS
<u>09008724</u>	5952344	250	01/19/1998	TETRAHYDROISOQUINOLINE DERIVATIVES
08813523	<u>5962472</u>	150		USE OF 4-PHENYL-3,6-DIHYDRO-2H-PYRIDYL DERIVATIVES
08789351	Not Issued	164	01/27/1997	TETRAHYDROISOQUINOLINE DERIVATIVES

Inventor Search Completed: No Records to Display.

	Last Name	First Name
Search Another:	Mutel	Vincent
Inventor		Search

To go back use Back button on your browser toolbar.

Back to PALM | ASSIGNMENT | OASIS | Home page